

### **Amendments To The Claims**

The listing of claims will replace all prior versions, and listings, of the claims in the application.

### **Listing Of Claims:**

Claims 1-15 (Cancelled)

Claim 16 (New) A method for producing needle-shaped crystals of Arg<sup>34</sup>GLP-1(7-37), said method comprising placing an aqueous solution of said Arg<sup>34</sup>GLP-1(7-37) at a temperature of between 20-25 degrees centigrade for a time sufficient to allow production of said crystals, wherein said aqueous solution has a pH of between 6 to 7 and comprises in addition to said Arg<sup>34</sup>GLP-1(7-37), 100-200 mM of an inorganic salt and 1-15% (w/w) ethanol.

Claim 17 (New) The method of claim 16, wherein said aqueous solution contains 2-10 mg/ml of Arg<sup>34</sup>GLP-1(7-37).

Claim 18 (New) The method of claim 17, wherein said aqueous solution comprises between 5-10 % ethanol (vol/vol).

Claim 19 (New) The method of claim 18, wherein said aqueous solution further comprises a buffer.

Claim 20 (New) The method of claim 19, wherein said buffer is bis-Tris.

Claim 21 (New) The method of claim 20 wherein the concentration of said buffer is between 5-10 mM.

Claim 22 (New) The method of claim 21, wherein said inorganic salt is NaCl.

Claim 23 (New) The method of claim 22 wherein the pH of said aqueous solution is between 6.2 and 6.6.

Claim 24 (New) The method of claim 19, wherein said pH of said aqueous solution is between 6.2 and 6.6.

Claim 25 (New) The method of claim 24, wherein said inorganic salt is NaCl.

Claim 26 (New) A method for producing an acylated glucagon-like peptide 1 (GLP-1) analogue, said method comprising:

- (a) placing an aqueous solution of said GLP-1 analogue at a temperature of between 20-25 degrees centigrade for a time sufficient to allow production of needle-shaped crystals of said GLP-1 analogue, wherein said GLP-1 analogue in said aqueous solution is Arg<sup>34</sup>GLP-1(7-37) and said aqueous solution has a pH of between 6 to 7 and comprises in addition to said Arg<sup>34</sup>GLP-1(7-37), 100-200 mM of an inorganic salt and 1-15% (vol/vol) ethanol; and
- (b) acylating the Arg<sup>34</sup>GLP-1(7-37) that was crystallized in step a).

Claim 27 (New) The method of claim 26, wherein the Arg<sup>34</sup>GLP-1(7-37) in the aqueous solution of step a) was recombinantly expressed in yeast.

Claim 28 (New) The method of claim 26, wherein prior to step a) the GLP-1 analogue Arg<sup>34</sup>GLP-1(7-37) is precipitated at a pH of about 5.4.

Claim 29 (New) The method of claim 26, wherein said aqueous solution in step a) contains 2-10 mg/ml of Arg<sup>34</sup>GLP-1(7-37).

Claim 30 (New) The method of claim 29, wherein said aqueous solution in step a) comprises between 5-10 % ethanol (vol/vol).

Claim 31 (New) The method of claim 30, wherein said aqueous solution in step a) further comprises a buffer.

Claim 32 (New) The method of claim 31, wherein said buffer is bis-Tris.

Claim 33 (New) The method of claim 32 wherein the concentration of said buffer is between 5-10 mM.

Claim 34 (New) The method of claim 33 wherein said inorganic salt in step a) is NaCl.

Claim 35 (New) The method of claim 34 wherein the pH of said aqueous solution in step a) is between 6.2 and 6.6.

Claim 36 (New) The method of claim 31, wherein said pH of said aqueous solution is between 6.2 and 6.6.

Claim 37 (New) The method of claim 36, wherein said inorganic salt is NaCl.

Claim 38 (New) A method for producing crystals of exendin-4, said method comprising placing an aqueous solution of said exendin-4 at a temperature of between 4-37 degrees centigrade for a time sufficient to allow production of said crystals, wherein said aqueous solution has a pH of  $pI < pH < pI + 2$  and comprises in addition to said exendin-4, at least 25mM of a salt and at least 0.5% (vol/vol) organic solvent.

Claim 39 (New) The method of claim 38, wherein said aqueous solution contains from 0.5 to 20 mg/ml of exendin-4.

Claim 40 (New) The method of claim 38, wherein said aqueous solution contains from 2-10 mg/ml of exendin-4.

Claim 41 (New) The method of claim 38, wherein said salt is an inorganic salt.

Claim 42 (New) The method of claim 41, wherein said inorganic salt is present in said aqueous solution in a concentration of 100-200 mM.

Claim 43 (New) The method of claim 38 wherein said aqueous solution comprises 1-15% (vol/vol) of organic solvent.

Claim 44 (New) The method of claim 38, wherein said aqueous solution has a pH of  $pI < pH < pI + 2$ .

Claim 45 (New) The method of claim 38, wherein said aqueous solution further comprises a buffer.

Claim 46 (New) The method of claim 38, wherein said aqueous solution is placed at a temperature of between 20-25 degrees centigrade for a time sufficient to allow production of said crystals.

Claim 47 (New) The method of claim 40, wherein said aqueous solution is placed at a temperature of between 20-25 degrees centigrade for a time sufficient to allow production of said crystals.

Claim 48 (New) The method of claim 47, wherein said salt is an inorganic salt.

Claim 49 (New) The method of claim 48, wherein said inorganic salt is present in said aqueous solution in a concentration of 100-200 mM.

Claim 50 (New) The method of claim 49 wherein said aqueous solution comprises 1-15% (vol/vol) of organic solvent.

Claim 51 (New) The method of claim 50, wherein said aqueous solution has a pH of  $pI < pH < pI + 2$ .

Claim 52 (New) The method of claim 51 wherein said aqueous solution further comprises a buffer.